Amendments to the Claims:

The following listing of claims will replace all prior versions, and listings, of claims in the application:

- 1. (Currently Amended) A-An injectable pharmaceutical composition containing 8 to 16 mg/mL of glycyrrhizin, 3 to 6 mg/mL of cysteine and 80 to 160 mg/mL of aminoacetic acid wherein substantially no sulfite is contained in the pharmaceutical composition.
 - 2. (Canceled)
- 3. (Currently Amended) The <u>injectable pharmaceutical composition according to claim 1</u>, wherein glycyrrhizin is monoammonium glycyrrhizinate.
- 4. (Currently Amended) The <u>injectable pharmaceutical</u> composition according to claim 1, wherein cysteine is cysteine hydrochloride.
 - 5-8. (Canceled)
- 9. (Currently Amended) The <u>injectable</u> pharmaceutical composition according to claim 1, wherein the concentration of cysteine is more than 70% after the composition is stored at 60°C for 14 days.
- 10. (Currently Amended) A-An injectable pharmaceutical composition containing 8 to 16 mg/mL (as glycyrrhizin) of monoammonium glycyrrhizinate, 4 to 8 mg/mL of cysteine hydrochloride and 80 to 160 mg/mL of aminoacetic acid, wherein substantially no sulfite is contained in the pharmaceutical composition.
- 11. (Withdrawn-Currently Amended) A method of treating hepatic diseases comprising: administering intravenously to a patient a-an injectable pharmaceutical composition containing 8 to 16 mg/mL of glycyrrhizin, 3 to 6 mg/mL of cysteine and 80 to 160 mg/mL of aminoacetic acid, wherein substantially no sulfite is contained in the pharmaceutical composition.

- 12. (Withdrawn-Currently Amended) A method of treating allergy comprising: administering intravenously to a patient a an injectable pharmaceutical composition containing 8 to 16 mg/mL of glycyrrhizin, 3 to 6 mg/mL of cysteine and 80 to 160 mg/mL of aminoacetic acid, wherein substantially no sulfite is contained in the pharmaceutical composition.
- 13. (Withdrawn-Currently Amended) A method of treating hepatic diseases comprising: administering intravenously to a patient a-an injectable pharmaceutical composition containing 8 to 16 mg/mL (as glycyrrhizin) of monoammonium glycyrrhizinate, 4 to 8 mg/mL of cysteine hydrochloride and 80 to 160 mg/mL of aminoacetic acid, wherein substantially no sulfite is contained in the pharmaceutical composition.
- 14. (Withdrawn-Currently Amended) A method of treating allergy comprising: administering intravenously to a patient a-an injectable pharmaceutical composition containing 8 to 16 mg/mL (as glycyrrhizin) of monoammonium glycyrrhizinate, 4 to 8 mg/mL of cysteine hydrochloride and 80 to 160 mg/mL of aminoacetic acid, wherein substantially no sulfite is contained in the pharmaceutical composition.